A complex choreography of cell movements shapes the vertebrate eye

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SUMMARY

Optic cup morphogenesis (OCM) generates the basic structure of the vertebrate eye. Although it is commonly depicted as a series of epithelial sheet folding events, this does not represent an empirically supported model. Here, we combine four-dimensional imaging with custom cell tracking software and photoactivatable fluorophore labeling to determine the cellular dynamics underlying OCM in zebrafish. Although cell division contributes to growth, we find it dispensable for eye formation. OCM depends instead on a complex set of cell movements coordinated between the prospective neural retina, retinal pigmented epithelium (RPE) and lens. Optic vesicle evagination persists for longer than expected; cells move in a pinwheel pattern during optic vesicle elongation and retinal precursors involute around the rim of the invaginating optic cup. We identify unanticipated movements, particularly of central and peripheral retina, RPE and lens. From cell tracking data, we generate retina, RPE and lens subdomain fate maps, which reveal novel adjacencies that might determine corresponding developmental signaling events. Finally, we find that similar movements also occur during chick eye morphogenesis, suggesting that the underlying choreography is conserved among vertebrates.

KEY WORDS: Zebrafish, Eye, Morphogenesis, Cell tracking, Retina, Retinal pigmented epithelium (RPE), Lens, Fate map

INTRODUCTION

Organogenesis depends on an interplay between patterning and morphogenesis. In a dynamic tissue, such as the developing eye, cell movements can both control and be controlled by patterning. Despite extensive studies of developmental eye patterning (reviewed by Adler and Canto-Soler, 2007; Fuhrmann, 2010; Li et al., 2004; Martinez-Morales et al., 2004), the accompanying cell movements have remained largely unstudied.

Previous understanding of optic cup morphogenesis (OCM) comes from fixed tissue analysis (Hilfer, 1983; Hilfer et al., 1981; Schmitt and Dowling, 1994; Schook, 1980a; Schook, 1980b; Silver, 1981). The mature optic cup (OC) comprises neural retina (NR), retinal pigmented epithelium (RPE) and lens. OCM can be divided into four steps (Fig. 1A): evagination, elongation, invagination and rotation. Evagination initially generates a simple pouch on the side of the brain. The optic vesicle (OV) then elongates, and a furrow constricts the connection between eye and brain, forming the optic stalk. During invagination, the NR and RPE enwrap the lens as it emerges from the overlying ectoderm. Once the lens has pinched off from the ectoderm, OCM is considered to be complete. Shortly thereafter, postmitotic retinal neuron differentiation begins. Cellular mechanisms responsible for OCM have not been determined: although OCM is often depicted as a sequence of epithelial sheet folding events, this lacks any empirical underpinning.

Four cellular mechanisms determine tissue shape: cell division, death, shape change and movement. There is some knowledge of their roles in OCM. In chick, localized proliferation was controversially proposed to underlie certain steps (Hilfer et al., 1981; Schook, 1980a; Silver, 1981). In Xenopus, inhibiting proliferation leads to disorganized yet functional eyes, but the time point of OCM completion was not analyzed (Harris and Hartford, 1991); it is possible that disorganization arose during OCM rather than subsequently during retinal neurogenesis. In zebrafish, the role of proliferation has not been tested. Cell death has been reported to occur in spatiotemporally regulated patterns during amphibian, chick, mouse and rat OCM (Laemle et al., 1999; Martin-Partido et al., 2006; Rembold et al., 2006). Both studies identified novel cell death mechanisms behind ocular malformations, but little detailed analysis of OCM, or of RPE and lens.

More recently, in toto imaging and automated cell tracking were used to study cell movements underlying evagination (England et al., 2006; Rembold et al., 2006). Both studies identified novel cell movements and behaviors. OV cell movement is integrated with neighboring telencephalic and hypothalamic brain regions to coordinate forebrain neurulation and early evagination; mutant analysis revealed multiple mechanisms underlying cyclopia (England et al., 2006). Novel subdivisions of the early eye field behave in distinct manners to initiate OV evagination, with a first event being slowed midline convergence of future lateral OV cells (Rembold et al., 2006). Subsequently, it was shown that OV-specific downregulation of the cell adhesion molecule Ncad
mediates the slowed convergence (Brown et al., 2010). These highly informative studies, however, did not extend beyond initial eye morphogenesis stages.

Many signaling pathways have been implicated in specific OV patterning events, and mutations in these pathways can lead to morphogenetic defects. In zebrafish, FGF signaling patterns the anterior-posterior (AP) axis of the eye, and recent work revealed that patterning is integrated with the regulation of specific cell movements and epithelial cell cohesion (Picker and Brand, 2003; Picker et al., 2009). The zebrafish patched 2 mutant blowout exhibits upregulated Hedgehog pathway activity and coloboma, which is the defective closure of the choroid fissure (Lee et al., 2008). Mouse Hes1 knockouts exhibit failure of OC invagination and coloboma; this function during OCM may be independent of Notch signaling (Lee et al., 2005; Tomita et al., 1996). The mouse Lrp6 insertional mutant displays microphthalmia and coloboma, implicating the involvement of canonical Wnt signaling (Pinson et al., 2000; Zhou et al., 2008). Finally, inhibiting retinoic acid signaling in zebrafish and mouse also leads to invagination defects and coloboma (Lupo et al., 2011; Mic et al., 2004).

Cell-intrinsic mechanisms also regulate OCM: the medaka gene ojoplano appears to mediate basal constriction underlying invagination (Martinez-Morales et al., 2009). In addition, mouse ES cells grown under specific conditions can differentiate and self-organize into an OC structure without extraocular tissues (Eiraku et al., 2011). Clearly, however, intrinsic mechanisms must be coordinated with extrinsic signals in the embryo. Despite a growing body of work, we lack a comprehensive understanding of OCM and how morphogenetic defects arise.

We aimed here to gain a detailed understanding of the cellular events during vertebrate OCM. Does proliferation contribute to the basic morphogenetic program? When and where do cells move? Are movements temporally and spatially coordinated between retina, RPE and lens?

The optical accessibility of zebrafish embryos offered a unique opportunity to investigate OCM using 4D time-lapse imaging and cell tracking. We analyzed the contributions to OCM of cell division and cell movement, and mapped cell movements and generated fate maps for the component tissues of the eye. Our results identify novel morphogenetic events shaping the retina, RPE and lens, with important implications for their specification, and include studies of OCM in the chick embryo that indicate that this process is conserved across vertebrates.

**MATERIALS AND METHODS**

**Zebrafish**

Embryos (Tü or TL strains) were raised at 28.5-30°C and staged according to hours post-fertilization (hpf) and morphology (Kimmel et al., 1995).

**RNA synthesis and injections**

Capped RNA was synthesized using pCS2 templates (pCS2-EGFP-CAAX, pCS2FA-H2A.F/Z-mCherry, pCS2FA-Kaeade, pCS2FA-nls-Kaeade) and the Message mMachine SP6 Kit (Ambion), purified (Qiagen RNeasy Mini Kit) and ethanol precipitated; 300-500 pg RNA was injected into one-cell embryos.

**Cell cycle inhibition**

Embryos (10.5 hpf) were dechorionated and incubated in E3 (untreated), DMSO control (E3 containing 3% DMSO) or HUA (E3 containing 20 mM hydroxyurea, 150 μM aphidicolin, 3% DMSO) (Lyons et al., 2005). Embryos (24 hpf) were fixed in 4% paraformaldehyde (PFA), permeabilized in TBST (TBS containing 0.1% Triton X-100), blocked in TBST containing 2% BSA and incubated with anti-phospho-histone H3 antibody (1:250; Millipore, 06-570), then goat anti-rabbit secondary antibody (Invitrogen, A-11008) co-incubated with 1 μM TO-PRO-3 iodide (Invitrogen, T3605). Embryos were cleared in 70% glycerol. Labeled cells were counted manually, one eye per embryo (ImageJ, NIH).

**Imaging**

Embryos (12 hpf) were dechorionated, embedded in 1.5% low-melting-point agarose (in E2 plus gentamycin) in DeltaT dishes (Biotecophs, #0420041500C). Images were acquired using an Olympus FX1000 laser-scanning confocal microscope. Plates were overlaid with E2 plus gentamycin and covered to prevent evaporation. No stage heater was used; the room was 27.6±0.4°C and the stage slightly warmer. Based on when the lens pinches off (24 hpf), we estimate that sample temperature was very close to 28.5°C. Four-dimensional datasets were acquired: 38 z-sections, 3.52 μm z-step, 3.15 minutes between z-stacks, 40× water-immersion objective (1.15 NA).

Kaede photoconversion used Olympus Fluoview software to expose a region of interest (ROI) to 405 nm light for 15-20 seconds. Efficiency was assayed by photocconversion of green to red fluorescence in the ROI.

**Image processing and analysis**

Before cell tracking, 4D datasets were processed: files were converted to multilayer single time point TIFFs; using the Float Morphology plug-in (ImageJ), the ‘open’ function was applied for the nuclear channel and the ‘close’ function for the membrane channel; the membrane channel was subtracted from the nuclear channel to improve nuclear separation; finally, intensity was adjusted along the z-axis to compensate for dimmer signals in deeper, ventral tissue.

Images were processed using ImageJ and Adobe Photoshop. Volume rendering was performed using Amira (Visage Imaging) or Fluorender (Wan et al., 2009). Three-dimensional cell speed was quantitated using Excel (Microsoft). Quantitative data were analyzed using analysis of variance (ANOVA) and P-values determined using Tukey’s HSD test.

**LongTracker**

For cell tracking, we used the MATLAB-based program LongTracker to select nuclei at specific time points and z positions, while simultaneously viewing and resolving and adjacent frames in z and t, stepping through the dataset forward or backward in time. For accurate cell tracking, we required at least 50% nuclear pixel overlap between time points. Data were exported as a spreadsheet of nuclear centroid position and nuclear TIFF images. Trajectories were checked for discontinuities in four dimensions using Fluorender.

**Volume measurements and cell counting**

Tissue volumes were measured using Amira after manual segmentation based on the membrane channel. Using the same manual segmentation, cell number was quantitated using the nuclear channel and the program EVE (T. Shimada, University of Tokyo), after applying an interpolation step to decrease the anisotropy of the data. Between each pair of adjacent slices a new slice— the average of the pair – was inserted.

**Visualization and quantitation of mitoses**

Mitoses were marked using LongTracker. Classification as a mitosis required unambiguous identification of a metaphase plate and two subsequent daughter nuclei. LongTracker allowed us to identify mitoses that spanned multiple z-slices while avoiding double counting. Mitosis tracking data were visualized in four dimensions (FluoRender) and quantitated (Excel).

**Chick fate mapping**

Fertilized White Leghorn chicken eggs were incubated at 38°C until the appropriate stage (Hamburger and Hamilton, 1992). Embryos were removed from eggs using filter paper rings and cultured, ventral side up, in dishes filled with agarose-albumen medium (0.2% agarose, 50% albumen) at 37°C, 10% CO₂ (EC culture) (Chapman et al., 2004). For Dil injection, embryos were flipped dorsal side up, and access to the OV attained via a small incision in the vitelline membrane. Dil (Molecular Probes, D282; 0.25% in 1:20 DMSO:tetraglycol) or CM-Dil (Molecular Probes, D1302) was injected into one-cell embryos.
Probes, C7000; 0.125% in 1:2 DMSO:tetraglycol) was applied; embryos were immediately imaged and returned to culture. Embryos were fixed with 4% PFA in PBS, counterstained with DAPI or Hoechst 33342, stepped through a glycerol/PBT (PBS containing 0.1% Triton X-100) gradient to a final concentration of 80% glycerol in double-distilled H2O, and reimaged. Imaging was performed with a Leica SP5 laser-scanning confocal microscope. Volume rendering and image analysis were performed using FluoRender.

RESULTS

Time-lapse imaging of eye morphogenesis

To visualize directly the cellular basis of OCM, we generated 4D datasets using time-lapsed fluorescence confocal microscopy. Embryos were labeled for membranes (EGFP-CAAX) and nuclei (H2A.F/Z-mCherry) by RNA injection at the one-cell stage and imaged 12-24 hours post-fertilization (hpf; supplementary material Movie 1; see Materials and methods). Since our datasets show little bleaching and no bright pyknotic nuclei indicative of phototoxicity, we believe that our imaging does not perturb OCM, except for a slight flattening (on the dorsal side) caused by mounting.

Single confocal images (Fig. 1B-F) from a representative dataset show membrane and nuclear labeling; volume renderings (Fig. 1B’-F’) illustrate changes in tissue shape and size. By 12 hpf (Fig. 1B-B’), the OV (purple) is a flat, wing-like structure emerging from the neural keel. By 14 hpf (Fig. 1C-C’), it has elongated posteriorly, assuming a flipper-like shape, and a furrow (Fig. 1A, arrowhead) drives anteriorly to constrict the connection between eye and brain, generating the presumptive optic stalk. OV invagination commences at 16 hpf (Fig. 1D-D’); the overlying ectoderm thickens, forming the lens placode (yellow). The lens is essentially round at 20 hpf (Fig. 1E-E’), as the OV enwraps it. Finally, at 24 hpf (Fig. 1F-F’), the lens has pinched off from the ectoderm and OCM is complete.

The optic vesicle increases in both volume and cell number during morphogenesis

As a first step toward determining underlying cellular mechanisms, we measured OV volume and cell number in 4D datasets. Its volume increased only moderately (46%) over 12 hours (from 0.84±0.06×10⁶ to 1.23±0.12×10⁶ μm³, mean+s.d.; n=3), with most of the increase between 12 and 16 hpf (Fig. 2A, supplementary material Fig. S1B). In previous measurements of fixed samples (Li et al., 2000), OV volumes were ~40% smaller due to fixation, but there was little change between 16 and 24 hpf, in agreement with our data.

Distinct lateral (ll) and medial (ml) OV layers are first distinguishable at 16 hpf (Fig. 1D). The lateral layer is often considered presumptive NR and the medial layer presumptive RPE (but see below). At 16 hpf, the layers were similar in size (ll, 0.65±0.11×10⁶ μm³; ml, 0.52±0.08×10⁶ μm³; Fig. 2A, supplementary material Fig. S1B). From 16-24 hpf (during invagination), this changes dramatically: the lateral layer expands to greater than four times the volume of the thinning medial layer (ll, 1.00±0.10×10⁶ μm³; ml, 0.23±0.02×10⁶ μm³; Fig. 2A, supplementary material Fig. S1A). The lens is first visible at ~15 hpf as a slight thickening of the overlying ectoderm. It grows steadily until ~23 hpf, then pinches off from the ectoderm, comprising ~10% of the OC volume (0.13±0.02×10⁶ μm³; Fig. 2A, supplementary material Fig. S1A,B). Changes in relative volumes were highly reproducible (Fig. 2A, supplementary material Fig. S1A,B).

During OCM, cell number roughly doubled, from 1142±78 (12 hpf) to 2467±76 (24 hpf) (Fig. 2B, supplementary material Fig. S1C), with half of the increase occurring from 12-16 hpf (1142±78 to 1805±222), before invagination. Relative cell numbers were very consistent between embryos (supplementary material Fig. S1C). Cell density increased gradually (supplementary material Fig. S1D).

Fig. 1. Timecourse of optic cup morphogenesis in zebrafish. (A) Schematic of optic cup morphogenesis (OCM), dorsal view. Arrowhead, furrow constricting the optic stalk. (B-F) Single confocal slices from 4D dataset of right optic vesicle (OV) at 12-24 hpf showing EGFP-CAAX (membranes, green) and H2A.F/Z-mCherry (nuclei, magenta). Dorsal views. (B’-F’) Volume renderings. OV (purple), lens (beige), brain (gray). (B’-F’) Dorsal views. (B”-F”) Lateral views. Asterisk, choroid fissure. br, brain; ov, optic vesicle; ec, ectoderm; ml, medial layer; ll, lateral layer; le, lens; A, anterior; P, posterior; M, medial; L, lateral; D, dorsal; V, ventral. Scale bar: 50 μm.
Can proliferation alone account for this increase in cell number? Mitoses are easily recognizable in our 4D datasets; H2A.F/Z-mCherry marks the elongated metaphase plate. We manually counted all mitoses (supplementary material Fig. S1E,F; see Materials and methods) and did not observe any obvious patterns of mitosis in our datasets (supplementary material Movie 2). The number of dividing cells was constant throughout OCM, with a slight plateau after ~21 hpf (Fig. 2B). Comparing changes in cell number with the number of mitoses shows that proliferation does not account for the rapid increase in cell number from 12-16 hpf (417±90 mitoses versus 663±144 additional cells; Fig. 2B,C). This suggests that, surprisingly, cells continue to enter the OV after 12 hpf, when evagination is thought to be complete.

We therefore directly counted cells moving from brain to eye after 12 hpf, using a nuclear-localized (nls) version of the fluorophore Kaede (Fig. 2D-H), which is converted from green to red fluorescence by irreversible photocleavage (Ando et al., 2002). At 12 hpf, the entire OV was photoconverted (Fig. 2D,E). Thus, any green nuclei in the eye thereafter (omitting overlying ectoderm/lens) must have entered from the brain during extended evagination. The 235±64 cells moving from brain to OV between 12 and 16 hpf neatly account for the difference between the increase in cell count (663±144) and mitotic count (417±90) (Fig. 2C). Late evagination ends by 16 hpf; the fraction of green nuclei thereafter remained roughly constant (~19.1%; Fig. 2H). Interestingly, late-moving cells specifically populated the ventroanterior OC (Fig. 2F,G). A similar extended evagination, in which late-moving cells populate the ventral OC, was reported in Xenopus, although without a detailed timecourse (Holt, 1980). Thus, this phenomenon might be a conserved feature of vertebrate OCM.

After 16 hpf, the number of mitoses matched or exceeded the change in cell number (Fig. 2C), implying that cell division and death can completely account for changes in cell number. However, although these quantitative analyses reveal the role of proliferation in controlling cell number, they do not address its role in morphogenesis.

**Mitosis is dispensable for basic aspects of OCM**

Inhibiting proliferation does not affect retinal cell type differentiation in Xenopus; however, the resulting eyes are small and disorganized (Harris and Hartenstein, 1991). OCM completion, however, was not analyzed as a time point and it is a formal possibility that loss of mitosis specifically during OCM could have
caused disorganization. In addition, this has not been tested in zebrafish. To test the role of proliferation specifically during zebrafish OCM, we incubated embryos in the DNA synthesis inhibitors hydroxyurea and aphidicolin (HUA) from 10.5-24 hpf, then analyzed effects on OCM. HUA treatment, as assayed by phospho-histone H3 staining (Fig. 3D-F), inhibited mitosis by ~87% (untreated, 74±16 mitoses per eye; DMSO, 62±13; HUA treated, 8.3±3.8; Fig. 3G).

HUA-treated OVs contained significantly fewer cells (826±121) than untreated (2235±296) or DMSO-treated (1895±295) OVs (Fig. 3H). OV volume decreased modestly from 1.15±0.11×10^6 μm^3 (untreated) or 0.99±0.07×10^6 μm^3 (DMSO) to 0.84±0.12×10^6 μm^3 (HUA treated; Fig. 3I). HUA-treated cells were distinctly larger (Fig. 3B,C), as reflected in a roughly halved cell density [1.0±0.1 (HUA) versus 1.9±0.3 (DMSO) or 1.9±0.1 (untreated) nuclei/10^3 μm^3].

Despite effects on cell number and size, morphogenesis proceeded normally (Fig. 3A-C). HUA-treated eyes were smaller but not disorganized, containing recognizable NR, RPE and lens (Fig. 3C), suggesting that the previously observed disorganization arose during retinal neurogenesis. The retina enveloped the lens slightly less than normal, but this might be due to decreased retina and RPE cell number. The fraction of OC volume occupied by RPE was similar to that of controls (untreated, 0.21±0.02; DMSO treated, 0.21±0.05; HUA treated, 0.24±0.02; Fig. 3K), indicating that the RPE thins and the NR thickens normally. Therefore, although mitosis contributes to changes in cell number, organ volume and cell density, it is dispensable for the basic program of OCM.

Tracking cell movements during OCM

Setting mitosis aside, we hypothesized that OCM relies primarily on cell movements. Therefore, we tracked cell movements in four dimensions, mapping the discrete morphogenetic events underlying OCM. We developed a MATLAB-based program, LongTracker, for manual cell tracking (H.O., K.R.C., Y. Wan, K.M.K., C. Hansen, C. K. Rodesch and C.-B.C., unpublished; see Materials and methods). Accurate cell tracking allows us to analyze cell movements and interrogate 4D datasets for cell fates, either prospectively or retrospectively. Importantly, many cells can be tracked from a single dataset, obviating the need to superimpose positional information from many embryos (a necessarily noisy process).

Although laborious, manual cell tracking was successful. In total, we tracked 153 cells from three datasets, with most tracking from the primary dataset and the two others used for confirmation (supplementary material Table S1). In the primary dataset, ~3% of OC cells were tracked from retina, RPE and lens. All data shown in the following figures are derived from the primary dataset. Combining our cell tracking with ImageJ, Amira and FluoRender for visualization, we defined the component cell movements of OCM and the timeline of movements, and generated a fate map of eye subdomains based on cell position and trajectory.

Retinal cell movements

We defined four major prospective NR movements: extended evagination, elongation (pinwheel movement), rim movement and anterior rotation.

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**Fig. 3. Proliferation is dispensable for OCM.** (A-C) Confocal sections (live, 24-hpf) showing EGFP-CAAX (membranes, green) and H2A.FZ-mCherry (nuclei, magenta). (A) Untreated, (B) DMSO and (C) HUA embryos treated from 10.5-24 hpf. Insets show large HUA-treated cells versus DMSO control. (D-F) Confocal sections at 24 hpf showing phospho-histone H3 (green) and TO-PRO-3 (magenta) in (D) untreated, (E) DMSO-treated and (F) HUA-treated embryos. Dorsal view. (G-K) Quantitative analyses of (G) mitoses, (H) cell number, (I) OV volume, (J) cell density and (K) retinal pigmented epithelium (RPE) volume as a fraction of OC [neural retina (NR) plus RPE] at 24 hpf in untreated (U), DMSO-treated (D) and HUA-treated embryos. Numbers of embryos scored (one eye each) are indicated within each bar. *, P<0.05; **, P<0.01; error bars indicate s.d. Scale bars: 50 μm.
Fig. 4. Four major retinal cell movements during OCM. (A–D’) Extended evagination (12-16 hpf). Cells are moving from brain to eye. Arrowhead, embryo midline; dotted line, OV/brain boundary. (A–D) Trajectories over membrane channel average projection (grayscale). (A’–D’) Trajectories only. (E–G’) OV elongation (12-14 hpf). Thirteen cells (1-8 NR, 9-13 RPE) undergoing pinwheel movement. Arrowhead, leading edge of furrow. (E–G) Trajectories over membrane channel average (grayscale). (E’–G’) Trajectories only. (H, I) Kymographs showing change in AP position over time for NR (H) and RPE (I) cells. (J–M’) Rim movement (18-24 hpf). Cells are moving from the medial to lateral layer. (J–M) Trajectories over membrane channel average (grayscale). (J’–M’) Trajectories only. Dotted line (J) demarcates the medial and lateral layers. (N–Q’) Eye rotation (12-24 hpf). (N–Q) Trajectories over volume rendering of membrane channel (grayscale). (N’–Q’) Trajectories only. (R–T) Rotation summaries for (R) 12-16 hpf, (S) 16-23.9 hpf and (T) 12-23.9 hpf. White circles indicate origins. (A–G’, J–M’) Dorsal view. (N–T) Lateral view. Scale bars: 50 μm.
Zebrafish eye morphogenesis

Extended evagination (12-16 hpf)
Zebrafish OV evagination had been thought complete by 12 hpf (Schmitt and Dowling, 1994). Quantitative analysis shows that although most evagination is complete by 12 hpf, it continues for ~4 hours (Fig. 2D-H). Cell tracking also clearly demonstrates this (Fig. 4A-D′, supplementary material Movie 3). Four representative cells crossed from brain into OV, clearly occupying the OV by 17.2 hpf (Fig. 4C,D).

OV elongation: pinwheel movement (12-14 hpf)
OV elongation has been described in several vertebrates using fixed tissue analysis, but not its underlying mechanism (Hilfer, 1983; Schmitt and Dowling, 1994; Schook, 1980b; Silver, 1981). In zebrafish, the OV rapidly shifts from wing-like to a posteriorly elongated flipper-like shape (Fig. 1). In chick, localized proliferation was proposed to underlie OV elongation (Hilfer et al., 1981). We find only a slight posterior bias for mitosis during this period (58.7% of mitoses occur in the posterior OV; n=2). Instead, prospective NR and RPE cells (definitively identified by backtracking) participate in a rapid, coordinated pinwheel movement (Fig. 4E-G′, supplementary material Movie 4). Prospective NR streamed into the OV, moving posteriorly (tracked cells 1-3, 5) and laterally (cells 6-8) (Fig. 4H), while prospective RPE (cells 9-13) moved medially and anteriorly (Fig. 4I). Considering all tracked cells (not just the representatives shown), more cells moved posteriorly than anteriorly, resulting in net posterior growth. Underscoring the rapidity of this movement, prospective NR moved faster during elongation than during the following 2 hours (1.03±0.26 versus 0.84±0.14 μm/minute, mean±s.d.; supplementary material Fig. S2).

Rim movement (18-24 hpf)
The OV medial layer shrinks dramatically starting at 16 hpf, with concomitant growth of the lateral layer (Fig. 2A, supplementary material Fig. S1B). A previous 3D time-lapse study showed migration from the medial to lateral layer around the ventral rim, at the AP level of the optic stalk (Picker et al., 2009). We found such movements throughout the OV.

During invagination (18-24 hpf), medial layer cells involuted around the rim, joining the lateral layer, on both anterior and posterior margins of the OV (posterior shown, Fig. 4J-M′, supplementary material Movie 5). We saw no evidence of cells ingressing directly across the optic lumen, confirming a previous hypothesis (Li et al., 2000). Our data reveal the temporal extent of this movement (18-24 hpf) and, taken together with the ventral rim movement previously observed (Picker et al., 2009), show that rim movement occurs around most (if not all) of the OV circumference, providing an explanation for the reciprocal volume changes of the two layers.

Anterior rotation (12-24 hpf)
Based on fixed tissue analysis, it was proposed that the eye rotates during the two stages of cephalic flexure [14-15 hpf and 24-36 hpf (Schmitt and Dowling, 1994)], bringing the choroid fissure to its final ventral position. This rotation is clockwise in a right lateral view (counterclockwise in a left lateral view), distinct from the pinwheel movement described above and viewed in a perpendicular plane. At 24 hpf, the choroid fissure lies 20-30° anterior of the ventral pole of the eye (Fig. 1F′, asterisk).

Cell tracking revealed some rotation during the first stage of cephalic flexure (Fig. 4N-O′,R). However, this rotation continued over the next 8 hours, before the second stage (Fig. 4O-Q′,S), with all cells moving clockwise during OCM to different extents (Fig. 4T). Thus, eye rotation might not be a passive result of cephalic flexure, but an active independent motion. Alternately, cephalic flexure might not be restricted to distinct stages, but a continuous gradual movement.

Origin of retinal domains
Since we tracked cells throughout the NR and found little cell mixing, we could draw the first complete retina fate map. We clustered all tracked NR cells from the primary dataset (Fig. 5A,D, supplementary material Fig. S3A-F, Movie 6, Table S2) into dorsal, ventral, anterior, central and posterior domains, based on final 24-hpf position and trajectory shape (Fig. 5B,C,E,F, supplementary material Movie 7). Because not all cells were tracked, domain boundaries are somewhat imprecise. Within each domain, all cells traveled along similar trajectories (Fig. 5G-L, supplementary material Movie 8), except for ventral.

In a lateral view of the 12-hpf OV (Fig. 5C), the dorsal, anterior, ventral and posterior domains are arranged circumferentially, behind the laterally situated central domain. As OCM proceeds, the peripheral domains rotate circumferentially (clockwise, right lateral view), while the central domain inserts into the hub of the wheel (Fig. 5F). This model confirms and extends a previous anterior retina fate map (Picker et al., 2009).

A surprise was the central domain, which we separated out only after noting the unique, ‘gamma-shaped’ trajectories of the cells (Fig. 5I). Cells moved posteriorly during OV elongation, then executed a loop turn and moved medially during invagination. How does the central domain merge into the OC center? It might actively push into position, or instead might be passively encircled by other domains. To test both models, we tracked anterior, central and posterior cells (Fig. 5Q-U′, supplementary material Movie 9). From 14-18 hpf (Fig. 5R-S′), after elongation, central cells (blue) have not moved significantly, while posterior (yellow) and anterior (orange) cells have bypassed them on either side, effectively completing the merge. Subsequently, central cells finally begin to move medially as invagination ends (20-24 hpf; Fig. 5T-U′). We conclude that the central domain merges into the OC hub in a largely passive manner, while cells in other domains migrate around the central domain, encircling it.

To confirm these results independently, we performed Kaede fate mapping: small clusters of cells were marked and imaged at 12 hpf (Fig. 5M,O) and 24 hpf (Fig. 5N,P). Clusters stayed relatively compact, with occasional extraocular labeling due to photoconversion of overlying or underlying cells. We present two typical examples. Lateral OV cells approximately halfway along the AP axis at 12 hpf (Fig. 5M) resided in the central retina at 24 hpf (Fig. 5N), confirming central domain tracking. Medial OV cells (adjacent to the brain) approximately one-third from its anterior end at 12 hpf (Fig. 5O) resided in the anterior retina by 24 hpf (Fig. 5P). Extensive Kaede experiments (n=40) confirmed the NR fate map based on cell tracking.

RPE cell movements
Tracking revealed two major RPE movements: pinwheel movement and spreading.

OV elongation: pinwheel movement (12-14 hpf)
During OV elongation and anterior furrow progression, prospective RPE cells participated in the pinwheel movement described above. The representative RPE cells shown (tracked cells 9-13) moved in a single, coordinated pinwheel trajectory with NR (Fig. 4E-G′,
supplementary material Movie 4) medially and either slightly posteriorly (cell 9) or anteriorly with the furrow (cells 10-13; Fig. 4I). Again, cells moved rapidly, with a mean 3D speed of $1.17 \pm 0.17 \mu m/minute$ during elongation, slowing to $0.84 \pm 0.10 \mu m/minute$ in the subsequent 2 hours (supplementary material Fig. S2).

Spreading (14-24 hpf)
After pinwheel movement, the prospective RPE spreads dramatically to become a flat monolayer: the distance between nuclei increases greatly, and cells spread to cover the back of the NR (Fig. 6A-D', supplementary material Movie 10).

Fig. 5. Origin and movement of retinal subdomains. (A,D) Positions of tracked NR cells at 12 (A) and 24 (D) hpf. Dorsal views. (B,C) Volume rendering of retinal subdomains (12 hpf): posterior (yellow), central (blue), anterior (orange), dorsal (green) and ventral (red). (B) Dorsal view. (C) Lateral view. (E,F) Volume rendering of retinal subdomains (24 hpf). (E) Dorsal view. (F) Lateral view. (G-L) Retinal subdomain trajectories. Dorsal view. Circles indicate origins; arrowheads indicate termini. (G) Summary. (H-L) Individual subdomains. Insets illustrate typical trajectory shape, except for ventral, which has none. (M-P) Kaede fate mapping. Dorsal views, volume renderings. (M,O) OV at 12 hpf; photoactivated spot, magenta. (N,P) Final position after OCM (24 hpf). (M,N) Central domain. (O,P) Anterior domain. Dashed lines outline OV (12 hpf) and OC (24 hpf). (Q-U') Central domain merge (14-24 hpf): posterior (yellow), central (blue) and anterior (orange) trajectories. Dorsal views. (Q-U') Trajectories over membrane channel average (grayscale). (Q'-'U') Trajectories alone. nk, neural keel; br, brain; le, lens. Scale bars: 50 \mu m.
Origin of RPE domains

We generated an RPE subdomain fate map by clustering all tracked RPE cells in the primary dataset (Fig. 6E,H, supplementary material Fig. S3G-J, Movie 11, Table S2) into three domains – anterior, central and posterior – based on final 24-hpf position and trajectory shape (Fig. 6F,G,I,J, supplementary material Movie 12). Within each domain, cells traveled along similar trajectories (Fig. 6K, supplementary material Movie 8). Our fate map omitted the ventral eye, where diminished image quality precluded reliable tracking.

The posterior RPE domain initially abutted the posterior and dorsal retinal domains, whereas the anterior RPE initially abutted anterior and ventral retina. Interestingly, we find similar trajectories for neighboring RPE and retina cells. For example, posterior retina (Fig. 5G,H) and posterior RPE (Fig. 6K) both travel along a similar hook-shaped trajectory (supplementary material Movie 8). Therefore, neighboring tissues move in concert, regardless of eventual cell fate.

Again, we confirmed our manual tracking using Kaede fate mapping (n=7). A photoactivated spot within the anterior RPE domain at 12 hpf (Fig. 6L) gave rise to anterior RPE cells at 24 hpf (Fig. 6M, arrowhead) and some brain cells.

Lens cell movements

We observed two prospective lens movements: pinwheel movement and compaction.

OV elongation: pinwheel movement (12-14 hpf)

During OV elongation, surprisingly, prospective lens cells followed the same trajectory as underlying retinal cells (Fig. 7A-D, supplementary material Movie 13). Again, cells moved rapidly during elongation, with a mean 3D speed of 1.01±0.27 μm/minute, slowing to 0.80±0.12 μm/minute in the subsequent 2 hours (supplementary material Fig. S2). These speeds are strikingly similar to those of...
prospective retinal cells (1.03±0.26 μm/minute during elongation, slowing to 0.84±0.14 μm/minute). Therefore, although lens cells do not appear to directly contact the OV their movement appears coordinated, with the same direction and speed as underlying NR.

Compaction (12-24 hpf)
Prospective lens cells originate throughout the entire ectoderm overlying the OV. During lens formation, they compact dramatically. Skeletons were drawn linking seven anterior and seven posterior cells (Fig. 7E-H). From 12-24 hpf, the skeletons collapsed. Interestingly, although cells appeared to mix, the skeletons remained separated (see below).

Origin of lens domains
We generated a lens subdomain fate map. All tracked lens cells from the primary dataset (Fig. 7I,L, supplementary material Fig. S3K-L, Movie 14, Table S2) were clustered according to final 24-hpf position and trajectory shape. As noted above, we found more mixing among lens cells than among NR or RPE (Fig. 7E-H). Only one boundary was evident in this dataset, dividing anterior (superficial) from posterior (deep) lens halves at 24 hpf (Fig. 7M, supplementary material Movie 15). This boundary corresponded to a border dividing the AP axis of the ectoderm at 12 hpf (Fig. 7J,K, supplementary material Movie 15). The two skeletons (Fig. 7E-H) correspond to anterior and posterior lens, which appeared to be separated by a compartment boundary: although cells mixed within each domain, there was little or no mixing between them. Cells within each compartment traveled along similar trajectories (Fig. 7N, supplementary material Movie 8), with one exception: during pinwheel movement (12-14 hpf), the posteriormost lens cells stalled or moved anteriorly, initiating posterior lens compaction (data not shown).

Kaede fate mapping confirmed these results (n=17). Marking anterior ectoderm (12 hpf) yields the entire anterior lens (24 hpf), with few cells in the posterior half (Fig. 7O,P), which might be due
to imprecise photoactivation without a morphological landmark. Alternately, there might be a little mixing across the AP boundary that was overlooked in our tracking. Non-lens ectoderm remained labeled at 24 hpf (Fig. 7P), indicating that the prospective lens cells at 12 hpf are internixed with non-lens cells.

**Conservation of cell movements among vertebrates**

Having mapped cell movements underlying OCM in fish, we asked whether two surprising aspects were evolutionarily conserved: pinwheel movement underlying OV elongation, and the AP division of the lens ectoderm.

In chick, the primary OVs have formed by the seven-somite stage [ss; HH stage 9 (Hamburger and Hamilton, 1992)], and elongate through 19 ss (HH stage 13). The OV elongates laterally, rather than posteriorly as in zebrafish. To test for pinwheel movements, Dil (pseudocolored green) was applied to the OV at either the lateral margin (Fig. 8A) or anterior margin (Fig. 8C, arrowhead). During elongation, each lateral Dil mark (Fig. 8A) swept posteriorly and medially (Fig. 8B; n=12/12), while each anterior Dil mark (Fig. 8C, arrowhead) moved laterally (Fig. 8D; n=10/10). We conclude that cells are undergoing a stereotyped pinwheel movement like that in zebrafish (Fig. 8E).

Testing for pinwheel movement allowed us to simultaneously test for eye rotation during these early stages. Initially, the Dil labels were located on the lateral OV margin, distributed randomly along the dorsal-ventral axis (Fig. 8A’,C’). After elongation, lateral cells had moved dorsally (Fig. 8B’; n=9/11), while anterior cells had moved ventrally (Fig. 8D’, arrowhead; n=7/10). As with zebrafish, the extent of movement varied depending on position. Thus, the chick OV rotates (counterclockwise, left lateral view), as in zebrafish (Fig. 8F).

In chick, the lens placode forms at 20 ss (HH stage 13+) and has invaginated completely by 26-28 ss (HH stage 16). To test for conservation of the lens fate map, Dil was applied to anterior (Fig. 8G; n=9) or posterior (Fig. 8I; n=7) ectoderm overlying the OV prior to lens placode formation (8 or 10 ss). After OCM, anterior (Fig. 8G) and posterior (Fig. 8I) ectoderm gave rise to distinct lens domains (Fig. 8H,J). Occasionally, some labeling appeared to cross the boundary, probably because no morphological landmark guides initial labeling. We conclude that the lens fate map (Fig. 8K), with its AP boundary, is at least partly conserved between zebrafish and chick.

**DISCUSSION**

OC morphogenesis generates the fundamental structure of the eye. Here, to probe vertebrate OCM, we used 4D imaging and cell tracking, which present several advantages. First, live analysis eliminates fixation artifacts, yielding more accurate size and shape measurements. Second, tracking many cells from a single dataset allows us to precisely visualize movement of cells relative to one another and within tissue structure. Third, the high temporal resolution of the data allows us to easily discern changes in speed and trajectory. Finally, with these datasets collected, we can ask new questions in the future.

Cell tracking in four dimensions is a difficult computational problem: nuclei are often irregularly shaped, and cells can divide frequently and move quickly. Although advances have been made in automated 4D cell tracking (Bao et al., 2006; England et al., 2006; Fernandez et al., 2010; Keller et al., 2008; McMahon et al., 2006; Olivier et al., 2010; Rembold et al., 2006), in most cases it is not easy to apply existing algorithms to a given dataset. We therefore used manual tracking with our newly developed tool, LongTracker. Because of the limitations of

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**Fig. 8. Conservation of movements in chick OCM.** (A-F) Pinwheel movement and anterior rotation. Lateral (A,A’) or anterior (C,C’) OV was labeled and imaged immediately. (A,C) Ventral views. (A’,C’) Lateral views. (C) Two anterior Dil spots were marked: medial spot (asterisk), brain; lateral spot (arrowhead), OV. The lateral Dil spot moved medially (B) and dorsally (B’). The anterior Dil spot (arrowhead) moved laterally (D) and ventrally (D’). (B,D) Ventral views. (B’,D’) Lateral views. Dashed line outlines OV; dotted line indicates the midline. (E) Summary of pinwheel movement. (F) Summary of anterior rotation. (G-K) Lens fate map. Anterior (G) or posterior (I) ectoderm was labeled and imaged immediately. Ventral view. Dashed line outlines OV; dotted line indicates the midline. (H,J) The Dil label was found in distinct lens domains. Ventral view. Dashed line outlines OV; long-dashed line outlines lens. (K) Summary of lens fate map. Volume renderings: Dil and brightfield (A,A’,C,C’,G,G) or Dil and nuclear counterstain (B,B’,D,D’,H,J). Scale bars: 200 μm.
manual tracking, we tracked ~3% of OC cells; development of additional tools will facilitate in toto analysis (Megason and Fraser, 2003).

**Cellular basis of OCM**

Since proliferation is dispensable for OCM (Fig. 3), we focused on cell movements, mapping for the first time the movements that shape the vertebrate eye. During the first phase of OCM (12-14 hpf), the OV is still undergoing evagination; late-ingressing cells are destined for the ventral retina. Concurrently, all three tissues (NR, RPE, lens) undergo pinwheel movement, which results in OV elongation and optic stalk constriction. During the next phase (14-16 hpf), late evagination ends and cell movements shift: many cells (posterior NR, RPE, lens; central NR, RPE; dorsal NR) turn sharply. During the third phase (16-24 hpf), the OV invaginates: the RPE spreads, lens cells converge, and retinal precursors undergo rim movement, moving from the medial to lateral layer around most of the OV circumference. Concurrent with all of these movements, and spanning the entire 12-24 hpf period, the eye gradually rotates anteriorly in a plane perpendicular to that of pinwheel movement.

OCM is often depicted as a series of differently folded epithelial sheet models. We do not know the extent of epithelial folding here as we have not analyzed cell morphology or single cell behavior. However, previous work suggests that, in fish, single cell migration, rather than the movement of coherent epithelial sheets, drives OV evagination (Rembold et al., 2006). Further work is required to determine the extent to which non-epithelial cell behaviors drive subsequent steps of OCM.

Two previous studies have used in toto imaging to map cell movements during evagination stages (England et al., 2006; Rembold et al., 2006). Our work extends these analyses, beginning after the early evagination stages, by mapping cell movements through the subsequent stages of OCM. It will be interesting to integrate the data and determine whether regional behaviors, including subdomain boundaries, persist from neural plate stages through OC formation. The answers might be informative as to the signals regulating specific cell movements.

Although zebrafish OV evagination was described as complete by 12 hpf (Schmitt and Dowling, 1994), we find that it persists until ~16 hpf. Cells destined for the ventroanterior OC move from brain to eye after 12 hpf (Fig. 2C-F). A similar extended evagination in *Xenopus* (Holt, 1980) suggests evolutionary conservation. A failure to complete this movement might have implications for chorioretinal coloboma, which is characterized by faulty choroid fissure closure. Adding to the mechanisms that might underlie such a defect, we propose a novel one: if late ingressing cells (posterior) are not exposed to specific ectodermal signals, rendering them competent to propagate the initial wave of retinal ganglion cell differentiation. It was previously shown that initiation of *atoh5* (*atoh7*) expression occurs in tissue adjacent to the optic stalk (Masai et al., 2000); perhaps the juxtaposition of cells ingressing during late evagination (adjacent to the optic stalk) and central retina demarcates the initial wave of retinal ganglion cell differentiation.

Similar to chick, the zebrafish lens arises from a large ectodermal domain (Fig. 7) (Bhattacharyya et al., 2004). Surprisingly, this domain is subdivided into discrete regions, giving rise to lens domains that comprise functionally distinct cell types (Greiling and Clark, 2009; Lovicu and McAvoy, 2005). We hypothesize that anterior and posterior pre-lens ectoderm are exposed to different signals that initiate distinct differentiation pathways.

Backtracking cell movements allowed us to draw prospective fate boundaries earlier than any known gene expression pattern. For example, we fate mapped the RPE to a small domain at 12 hpf, whereas the first genes specific to the zebrafish RPE, *mitfa* and *mitfβ*, are not expressed until ~17 hpf (Lister et al., 2001). Combining gene expression data with our fate maps might suggest the signaling events involved in patterning the OV.

**Evolutionary conservation of cell movements**

To assess evolutionary conservation, we asked whether chick OV evagination involves pinwheel movements and anterior rotation and if the lens fate map exhibits an AP boundary. As all three answers were affirmative, we propose that the cell movements described here apply generally to vertebrate OCM.

However, there are differences between the chick and fish fate maps. For example, the chick OV posterior margin contributes to ventral OC, whereas in zebrafish it contributes to posterior and...
dorsal OC (Fig. 5) (Dutting and Thanos, 1995). How do we reconcile similar morphogenetic movements with fate map differences? Perhaps the same movements are used but to different extents. For example, pinwheel movement appears more dramatic in zebrafish than in chick; consequently, equivalent cells will be adjacent to different tissues and will subsequently encounter different inductive signals. Additionally, patterning signals might act differently between species. For example, FGF signaling in chick and mouse influences retina versus RPE fate (reviewed by Chow and Lang, 2001; Fuhrmann, 2010; Martinez-Morales et al., 2004), whereas in zebrafish it functions in AP retina patterning (Picker and Brand, 2005; Picker et al., 2009).

Conclusions
We combined 4D imaging and custom software to determine the cellular basis of vertebrate eye morphogenesis. This approach can be applied to other morphogenesis events, particularly in zebrafish as a favorable system for 4D imaging. To this end, we plan to make our LongTracker software freely available (H.O., K.R.C., Y. Wan, K.M.K., C. Hansen, C. K. Rodesch and C.-B.C., unpublished). The novel cell movements and fate maps described here represent a significant shift in our comprehension of OCM and should lead to improvements in molecular understanding and models of human eye disease.

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Competing interests statement
The authors declare no competing financial interests.

Supplementary material
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Supplementary material


